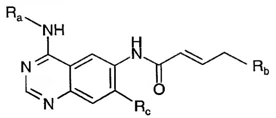
Abstract

A compound of general formula I



(I),

wherein:

R_a is a benzyl, 1-phenylethyl, or 3-chloro-4-fluorophenyl group;

R_b is a dimethylamino, *N*-methyl-*N*-ethylamino, diethylamino, *N*-methyl-*N*-isopropylamino, *N*-methyl-*N*-cyclopropylamino, *N*-methyl-*N*-(2-methoxyethyl)amino, *N*-ethyl-*N*--(2-methoxyethyl)amino, bis(2-methoxyethyl)amino, morpholino, *N*-methyl-*N*-(tetrahydrofuran-3-yl)amino, *N*-methyl-*N*-(tetrahydrofuran-2-ylmethyl)amino, *N*-methyl-*N*-(tetrahydrofuran-3-ylmethyl)amino, *N*-methyl-*N*-(tetrahydropyran-4-yl)amino, or *N*-methyl-*N*-(tetrahydropyran-4-ylmethyl)amino group; and

R_c is a cyclopropylmethoxy, cyclobutyloxy, cyclopentyloxy, tetrahydrofuran-3-yloxy, tetrahydrofuran-2-ylmethoxy, tetrahydrofuran-3-ylmethoxy, tetrahydropyran-4-yloxy, or tetrahydropyran-4-ylmethoxy group,

or a tautomer, stereoisomer, or salt thereof,

particularly the physiologically acceptable salts thereof with inorganic or organic acids or bases which have valuable pharmacological properties, in particular an inhibitory effect on signal transduction mediated by tyrosine kinases, their use in the treatment of diseases, especially tumoral diseases and diseases of the lungs and airways, and the preparation thereof.